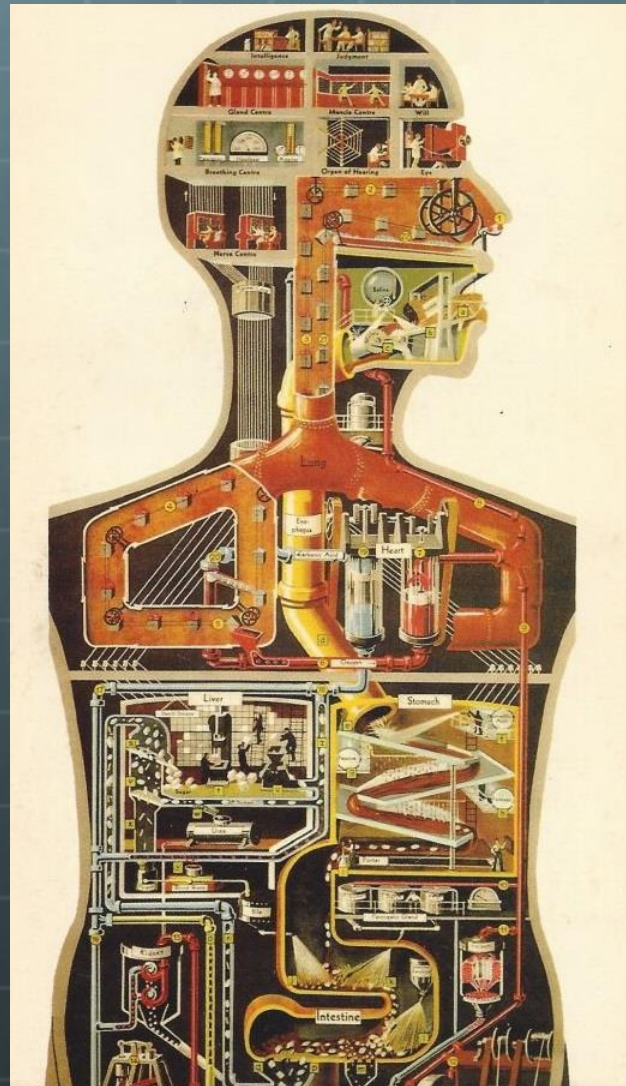


Synthetic Biology and iGEM



Genetically Engineered Machine (iGEM) Competition

Synthetic biology: what is it?

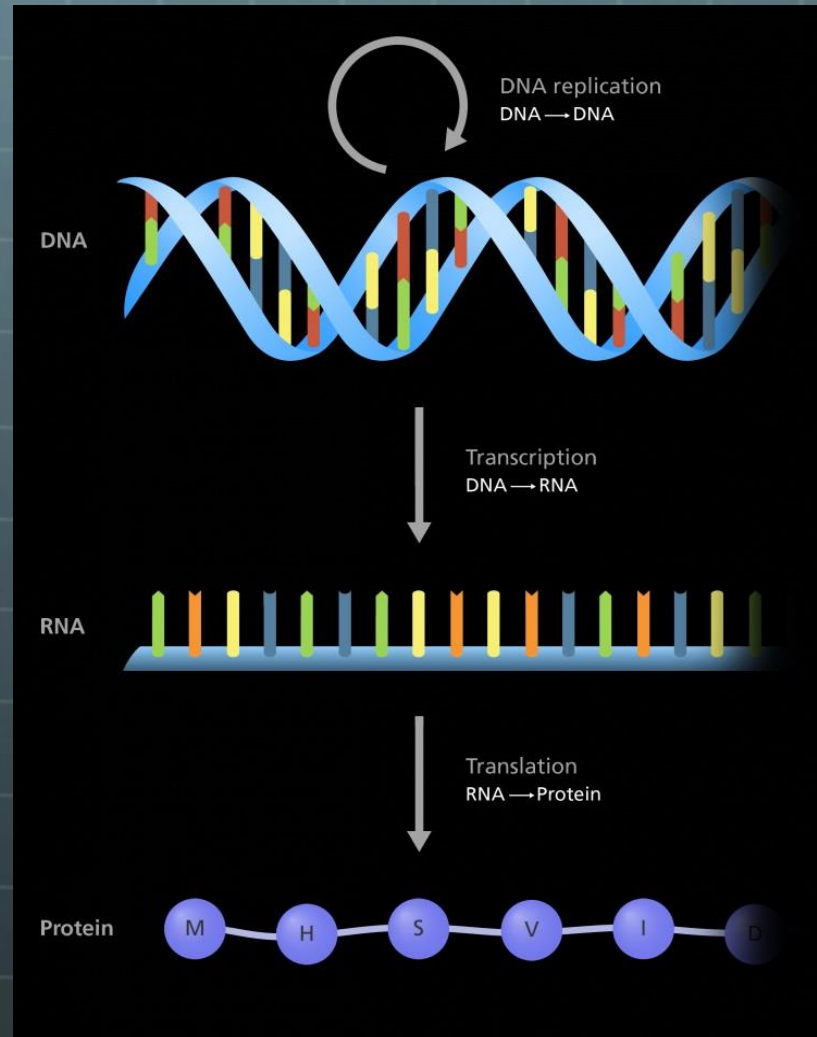


Cover of *Man Modified*, David Fishlock

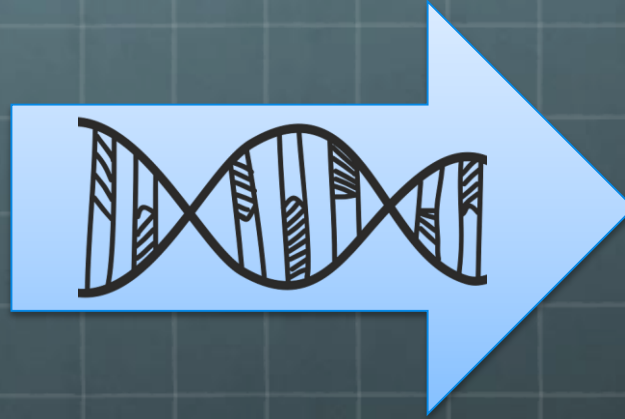
“The Lego of Life”



Transcription and Translation

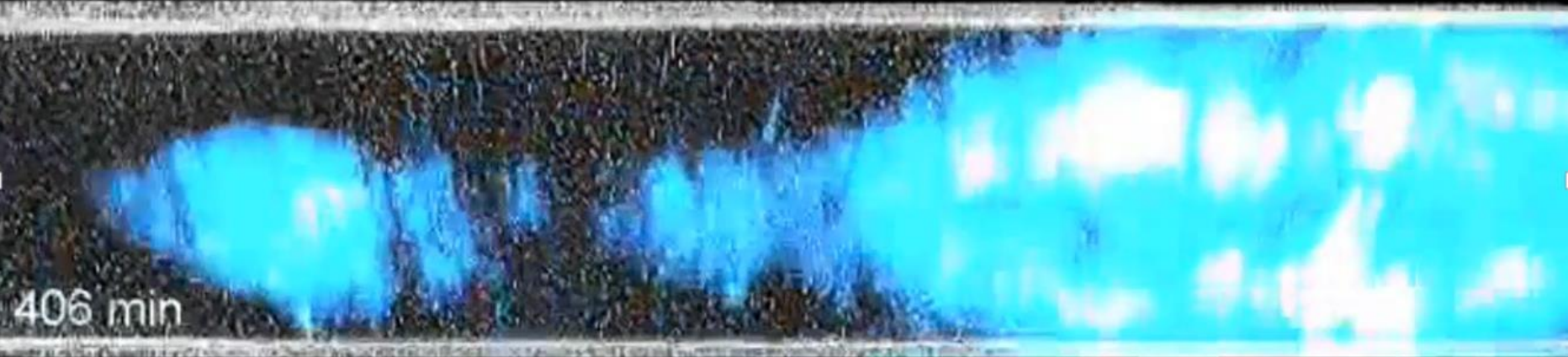


Synthetic biology: how do we do it?



naturevideo

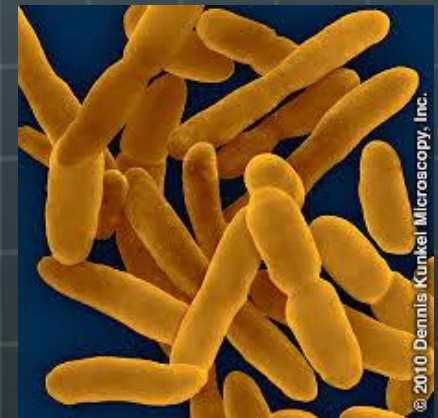
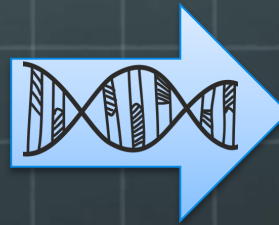
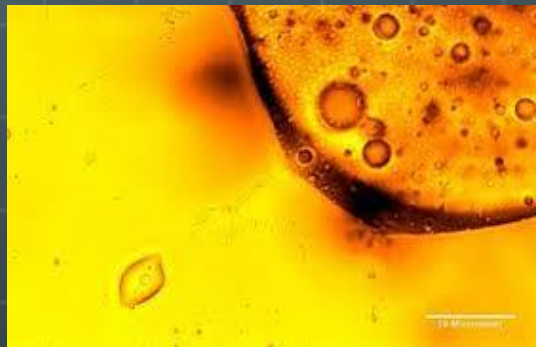
406 min



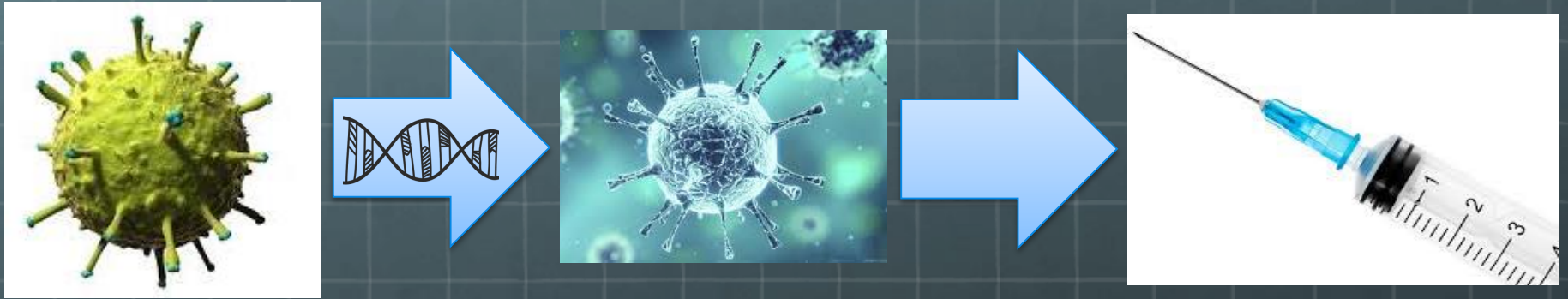
Synthetic biology: why is it important?



Synthetic biology: why is it important?



Synthetic biology: why is it important?



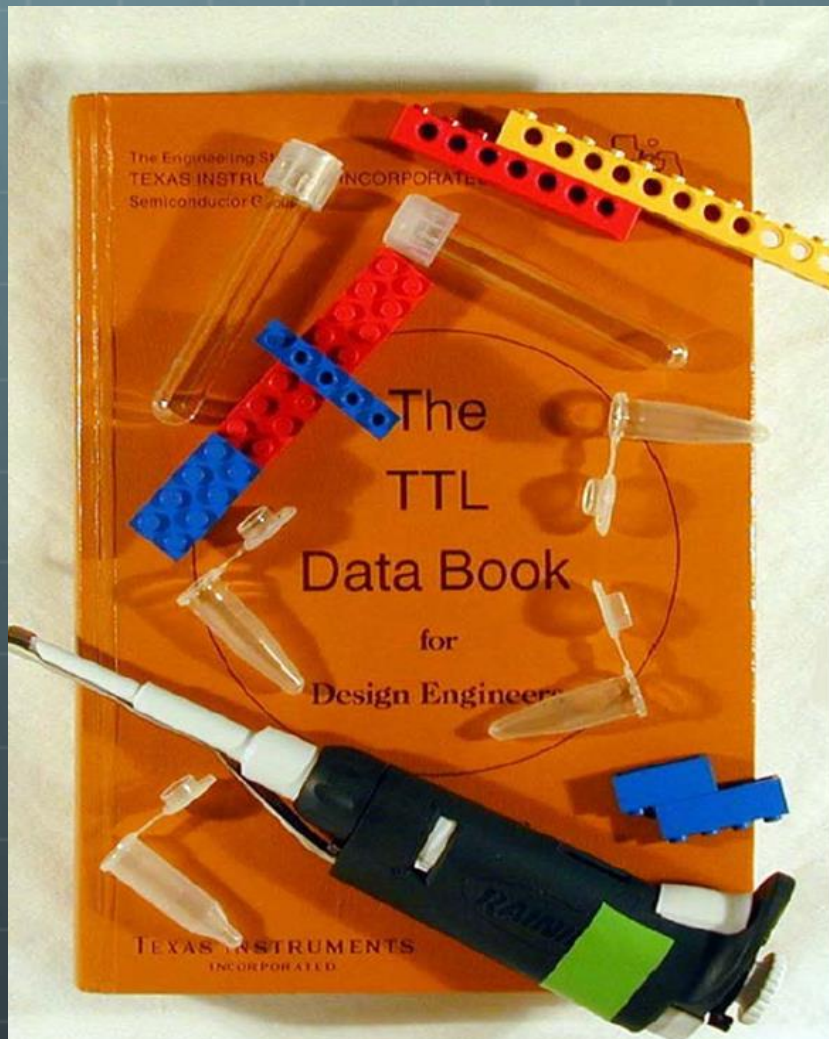


Genetically Engineered Machine (iGEM) Competition



iGEM Jamboree 2015

Registry of Standard Biological Parts



High copy number assembly plasmid backbones

The most common set of plasmid backbones that people use to assemble BioBrick® standard biological parts together are high copy BioBrick plasmid backbones. High copy plasmid DNA is easily purified in high yield from cultures, so it makes obtaining enough DNA for assembly easy.

The high copy plasmid backbones listed below have a common set of features.

1. A complete BioBrick® cloning site for easy cloning and assembly of BioBrick parts.
2. Terminators flanking the BioBrick® cloning site to insulate the vector from read-through transcription originating in the cloned BioBrick® part, device or system.
3. Primer binding sites for the standard BioBrick® verification primers VF2 (BBa_G00100) and VR (BBa_G00101). These primers are located for convenient sequencing and screening by colony PCR of cloned BioBrick® parts, devices, and systems.

Plasmid backbones are distributed by the Registry with a default insert. There are just a handful of default plasmid inserts used in the Registry. Many of the available plasmid backbones have the *ccdB* positive selection marker (BBa_P1010) as the default plasmid insert within the BioBrick® cloning site.

The *ccdB* gene ensures that when assembling two BioBrick® parts together, the uncut plasmid is not transformed. However, inclusion of the *ccdB* gene means that these vectors must be propagated in a *ccdB* tolerant strain, such as *E. coli* strain DB3.1 (BBa_Y1005).

Finally, to make assembly of BioBrick® parts easier, these BioBrick® assembly plasmid backbones are available with three different antibiotic resistance markers, so that you can use 3 antibiotic assembly methods to assemble BioBrick® parts.



-?-	Name	Description	Resistance	Replicon	Copy number	Chassis	Length
A/W	pSB1A3	High copy BioBrick assembly plasmid	A	pMB1	100-300		2157
A/W	pSB1A7	Transcriptionally insulated high copy BioBrick plasmid	A	pMB1	100-300		2431
A/W	pSB1AC3	High copy BioBrick assembly plasmid	AC	pMB1	100-300		3055
A/W	pSB1AK3	High copy BioBrick assembly plasmid	AK	pMB1	100-300		3189
A/W	pSB1AT3	High copy BioBrick assembly plasmid	AT	pMB1	100-300		3446
W	pSB1C3	High copy BioBrick assembly plasmid					2072
W	pSB1K3	High copy BioBrick assembly plasmid					2206
W	pSB1T3	High copy BioBrick assembly plasmid					2463



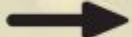
Karmella Haynes, an instructor of the 2006 Davidson College iGEM team, designed and constructed the plasmid backbone pSB1A7. You can read more about the 2006 Davidson project in their open-access paper *Engineering bacteria to solve the Burnt Pancake Problem* published in the *Journal of Biological Engineering*.



Robbie Bryant constructed the plasmid backbone pSB1AC3 in Tom Knight's lab.

Grand Prize Winner, 2015

UC Davis



Policy and
Practices

Pr
Eng

Olive Oil		Predicted Concentration (μM)		
Rancidity	Sample Number	$[S_M]$	$[S_L]$	$[U]$
Fresh	1	0.5	9.7	4.9
Fresh	2	1.8	7.7	5.6
Fresh	3	0.0	15.0	7.0
Fresh	4	0.0	20.7	6.4
Fresh	5	4.2	6.4	1.0
Fresh	6	0.0	15.0	1.8
Rancid	7	36.4	0.0	35.9
Rancid	8	104.6	0.0	28.3
Rancid	9	0.0	21.9	9.1



Saturated Short
Saturated Long
Unsaturated

nal Processing

Brainstorming

CRISPR

- soil enrichment
- fast into animals
- bacteria to control
- fertilizer to prevent algae blooms
- what is not
- established methods
- what animal cells
- taste?
- nutrients?

IGEM 2015 Project IDEAS

- Tar ball remediation by engineered bacteria or enzymes
- Biomass conversion to fuel (hydrocarbons)
- In vitro meat/ingredients
- engineer bacteria to make menthol, kill bad bacteria, mineralization?
- Sensors for water contaminants
- bacterial sensor
- paper
- colorimetric output
- toilet paper sensors
- data encryption in DNA
- parasites
- comprehensive + quick
- what to look for specifically

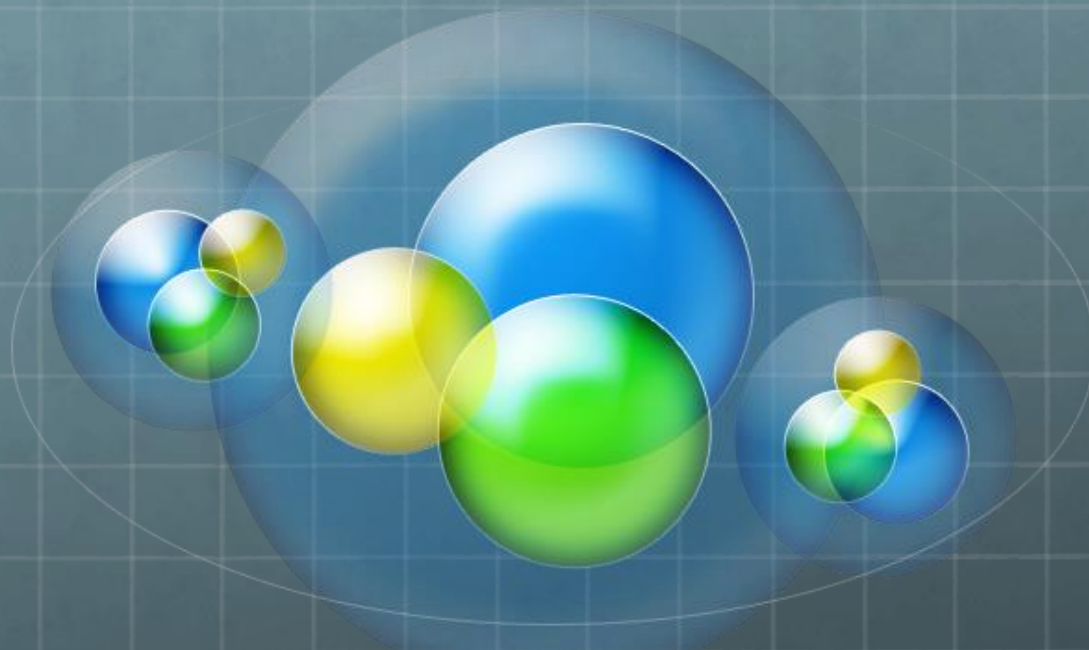
Other Ideas:

- decontaminating water
- plastic degrading bacteria
- follow upon engineered skin bacteria
- phage for antibiotic resistance
- THC testing
- HBV synthetic, other PEDs
- track glucose meter or pregnancy test
- archaeal mix for trash degradation using an engineered extremophile

Column/Bands
membrane/soluble chloride?

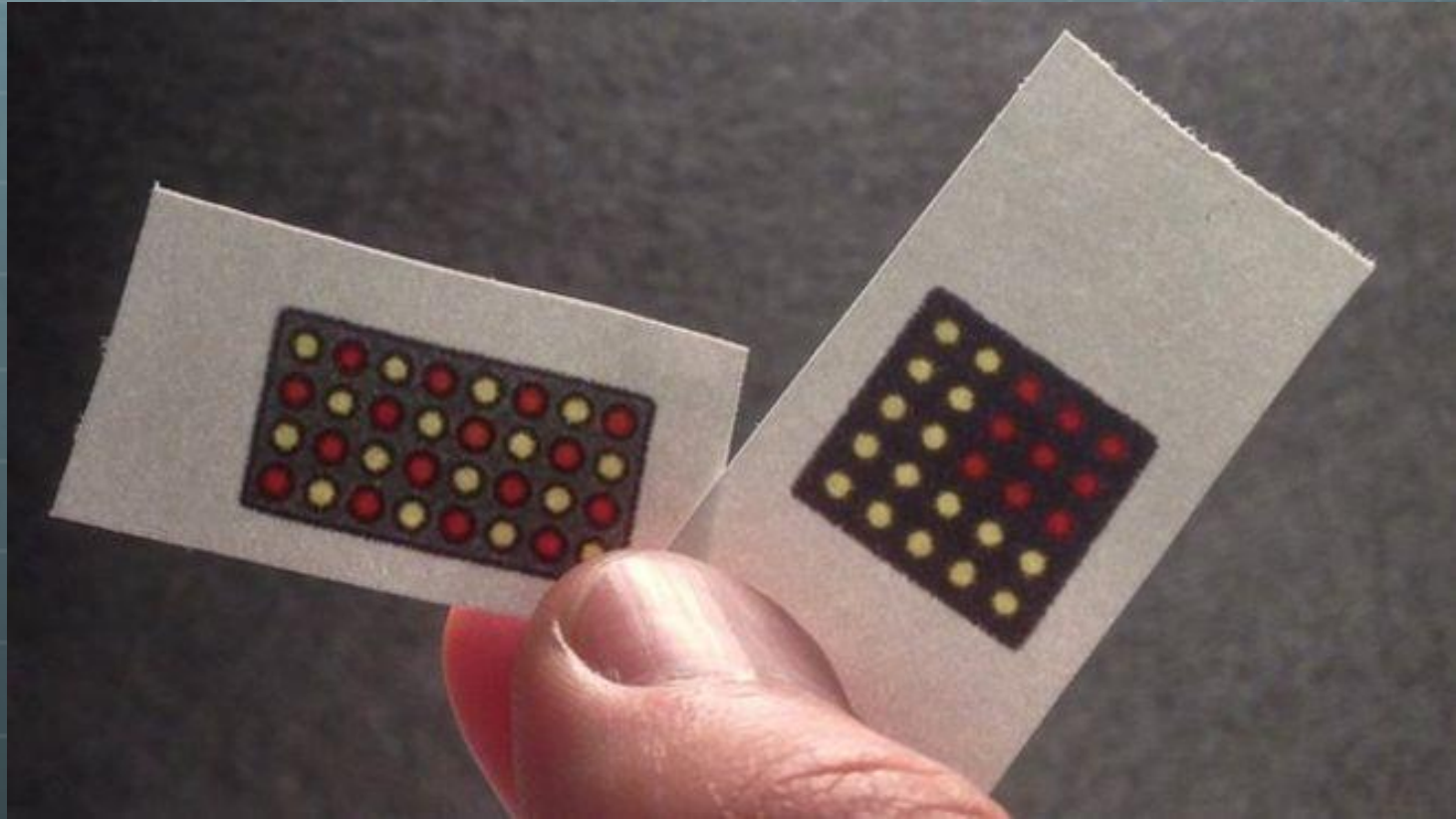
Other Notes:

- moving high-level pathogen from plant to bacteria (bacteriophage)
- what to make
- what to look for
- what to look for



Our Project: Paper-Based Cell-Free Thallium Sensor

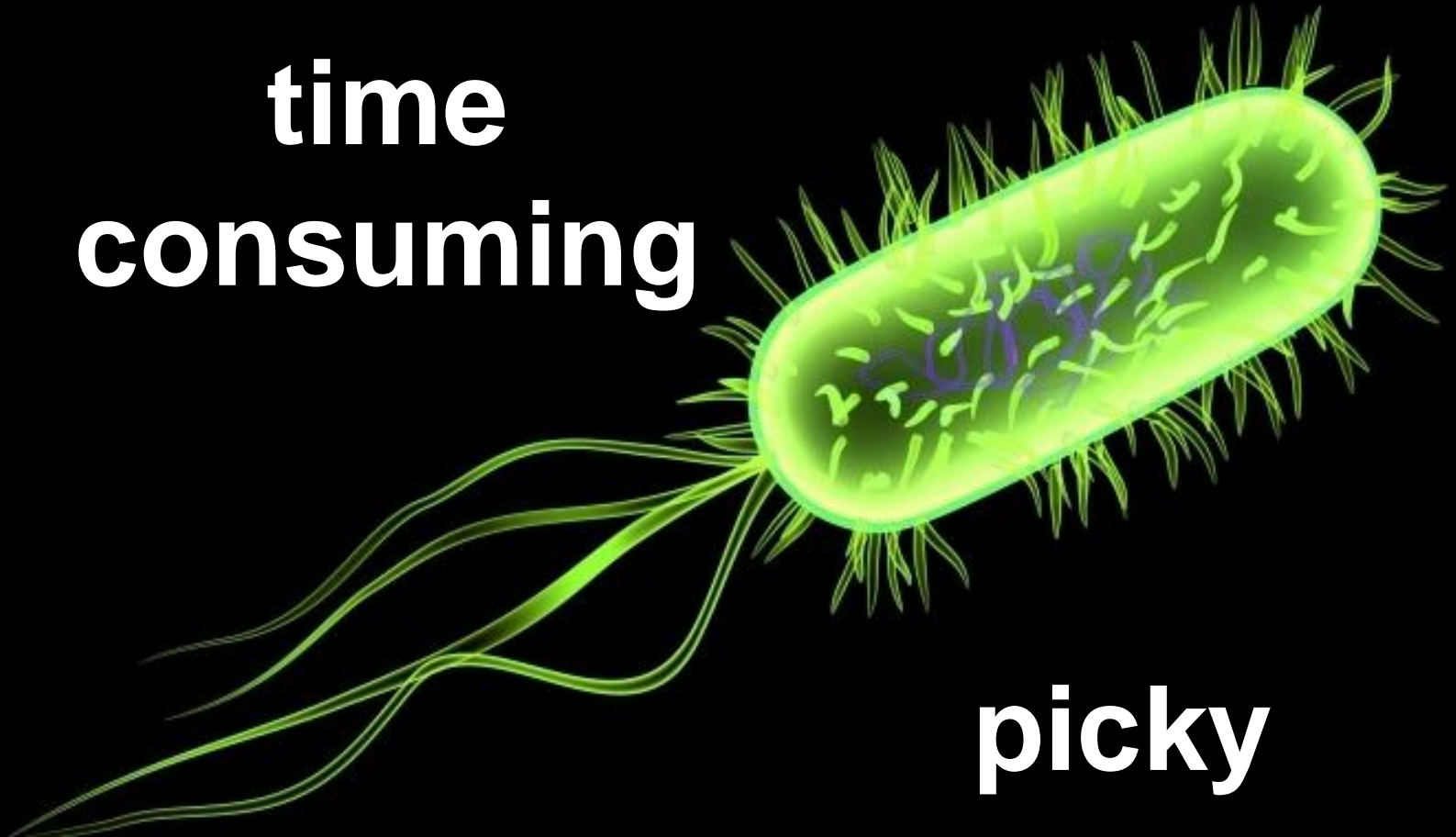
Paper-Based Sensor



BBC, "Prototype paper test can detect Ebola strains"

Cell-Free

**time
consuming**

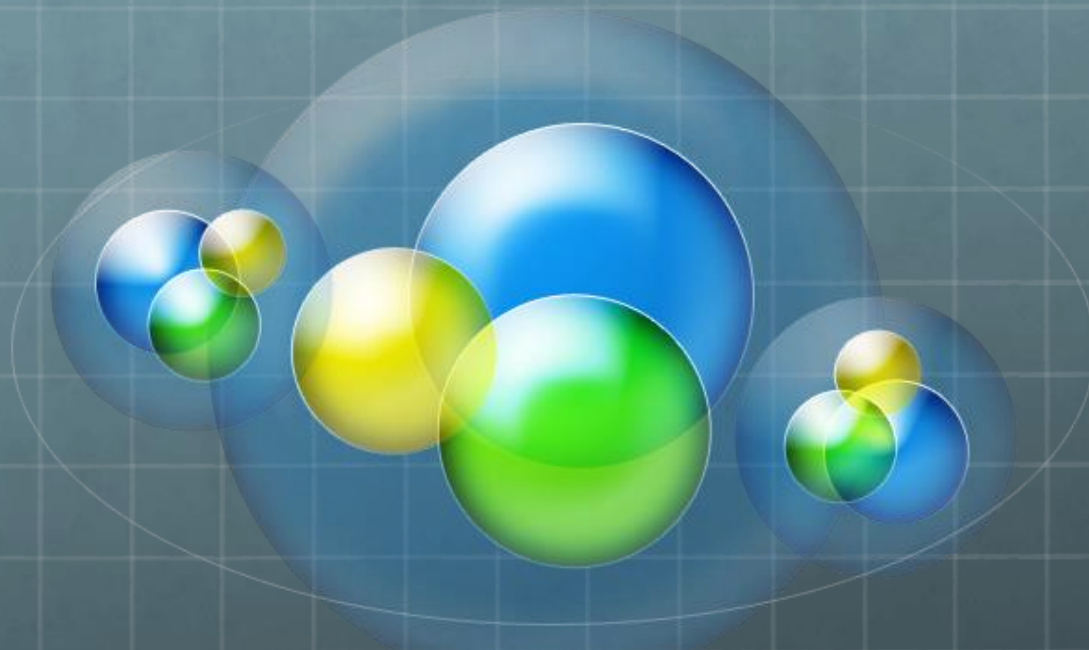


picky

Thallium

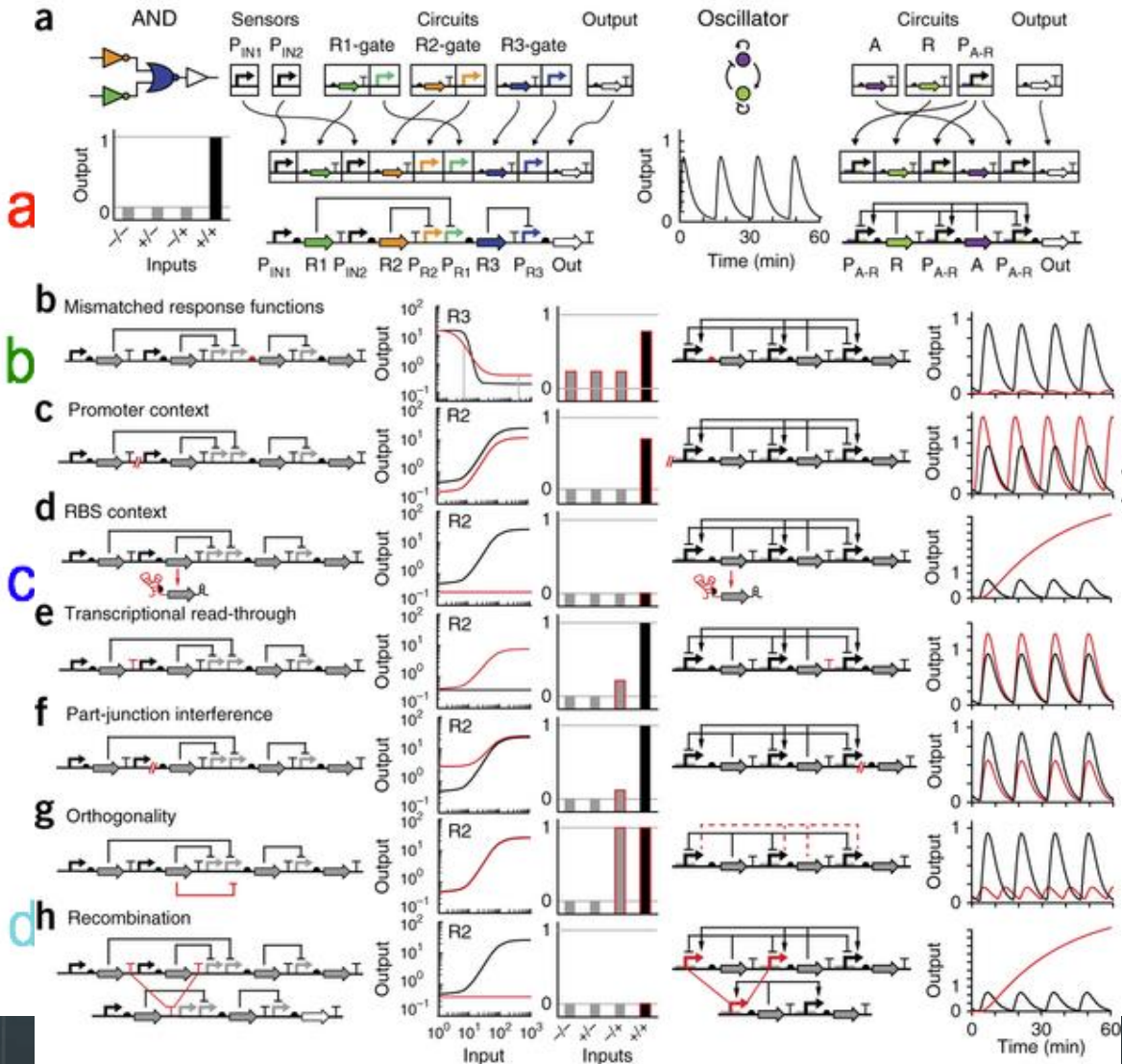
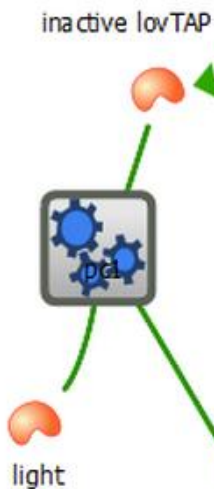
81
Tl
204.38



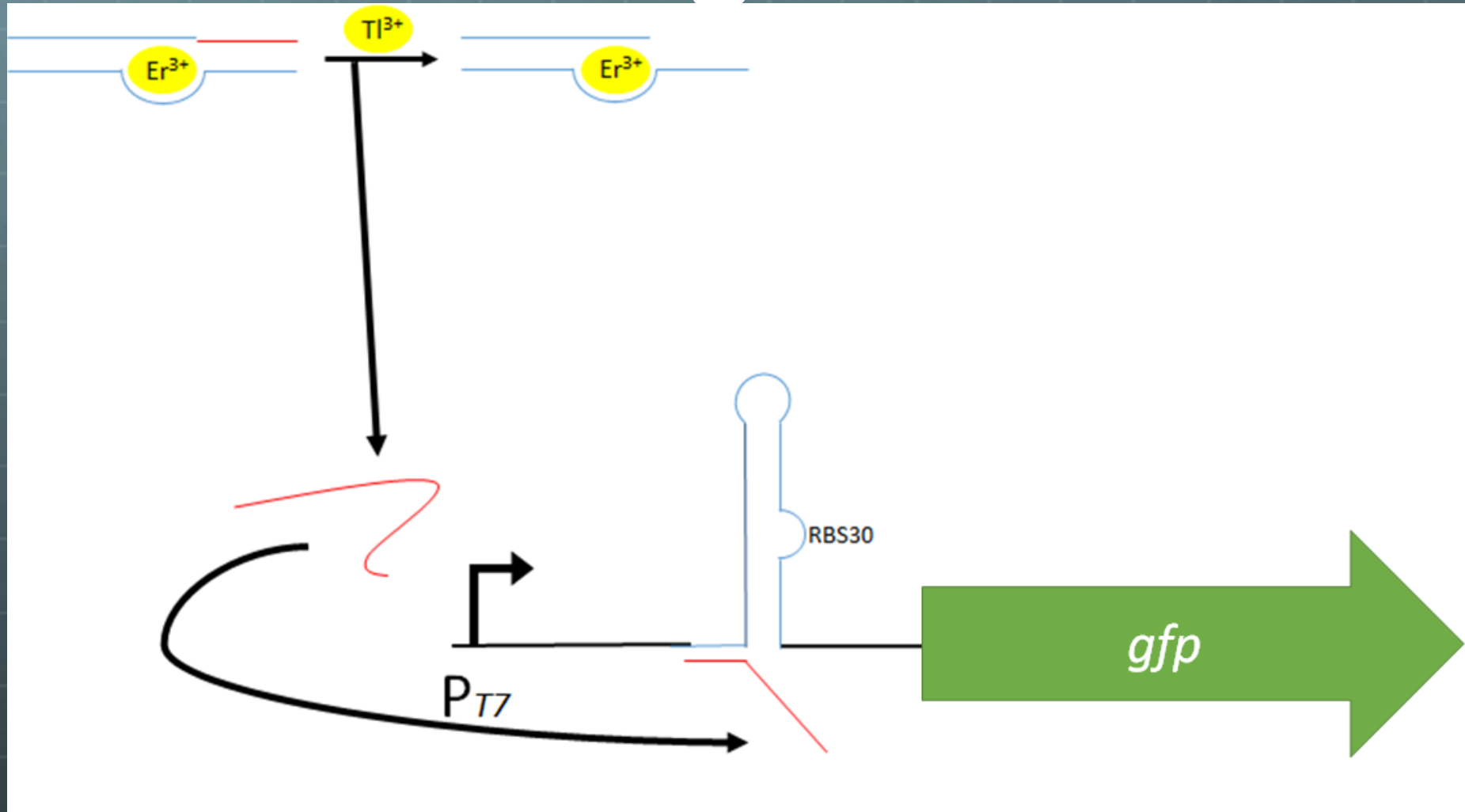


**Our Project:
Paper-Based Cell-Free Thallium
Sensor**

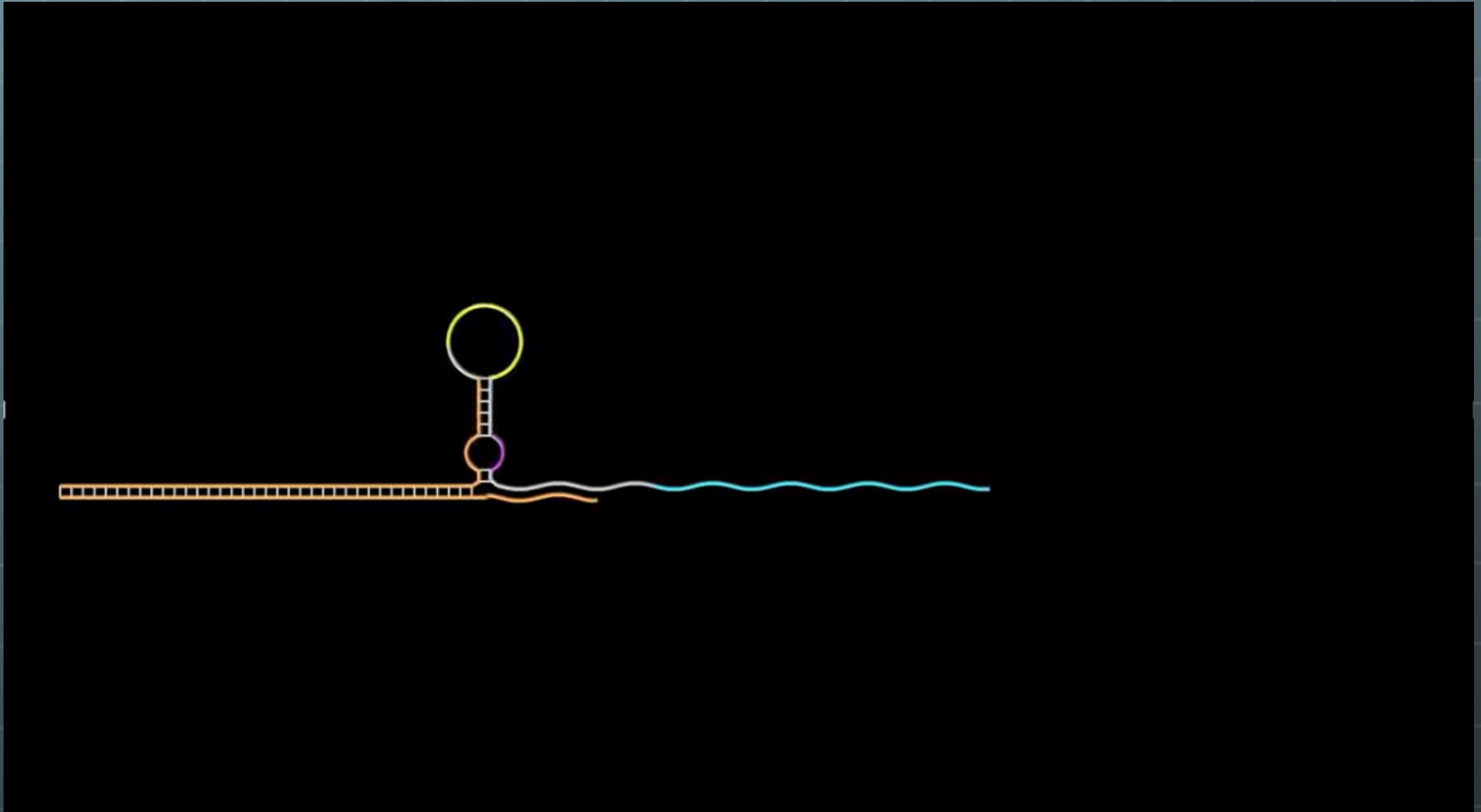
Genetic Circuits



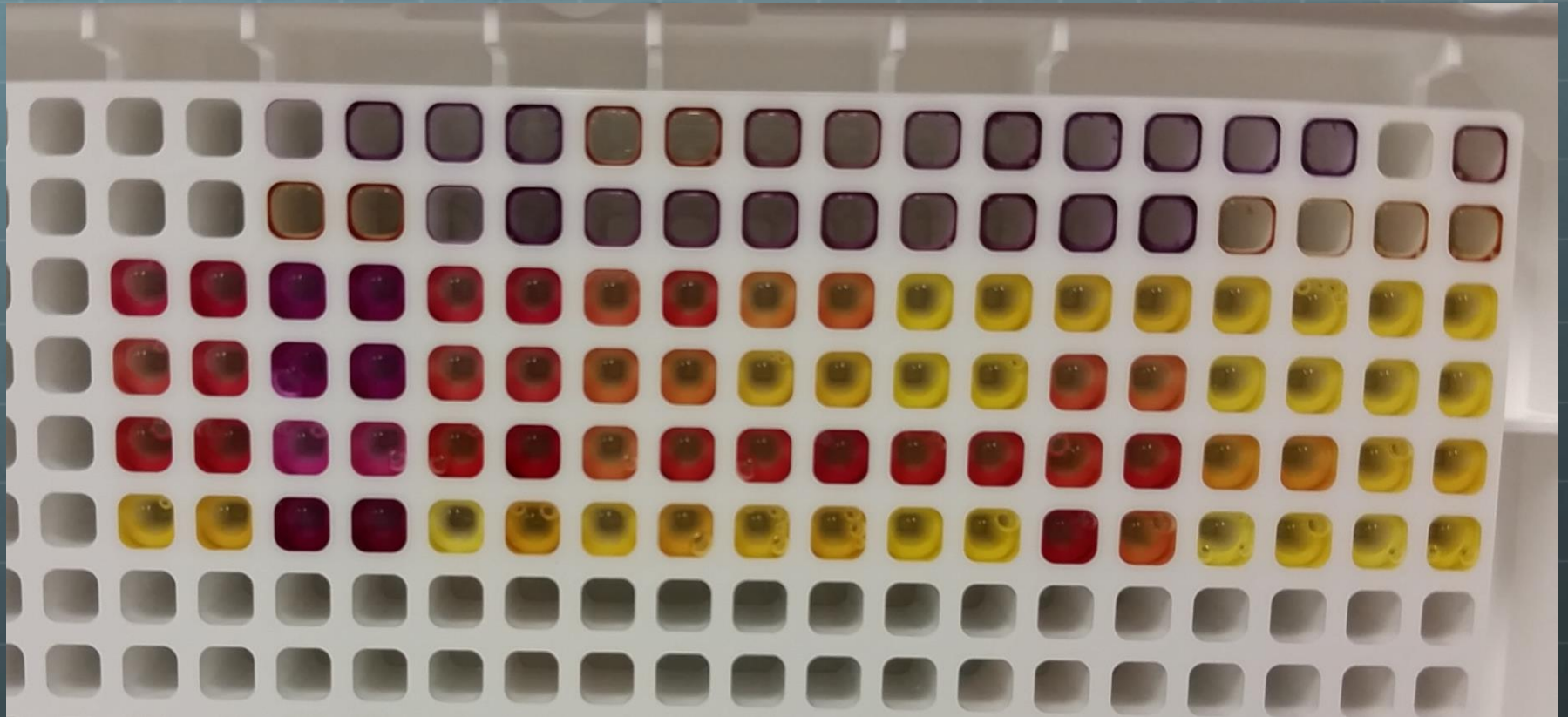
Genetic Circuit Diagram



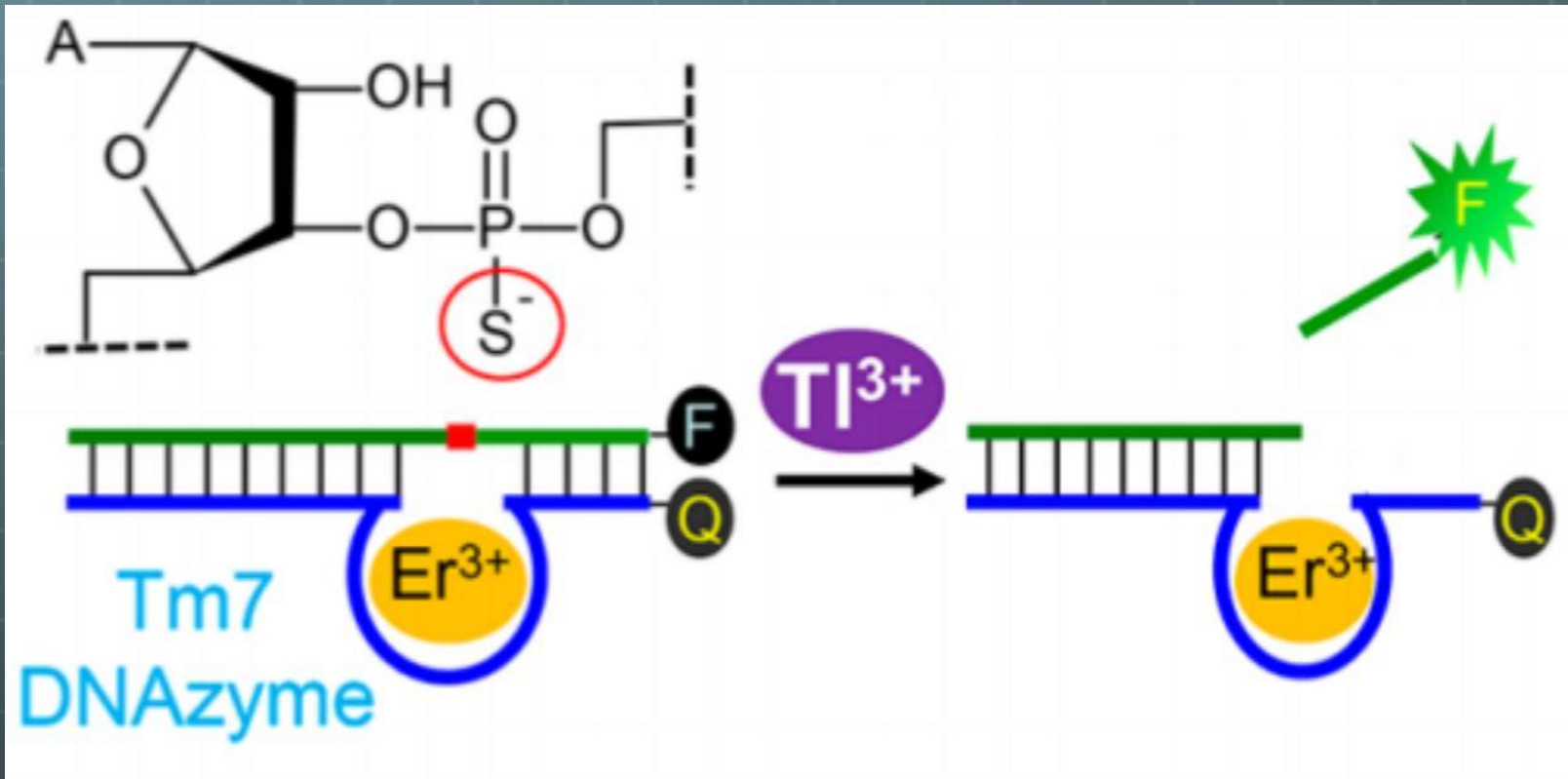
Toehold Switch



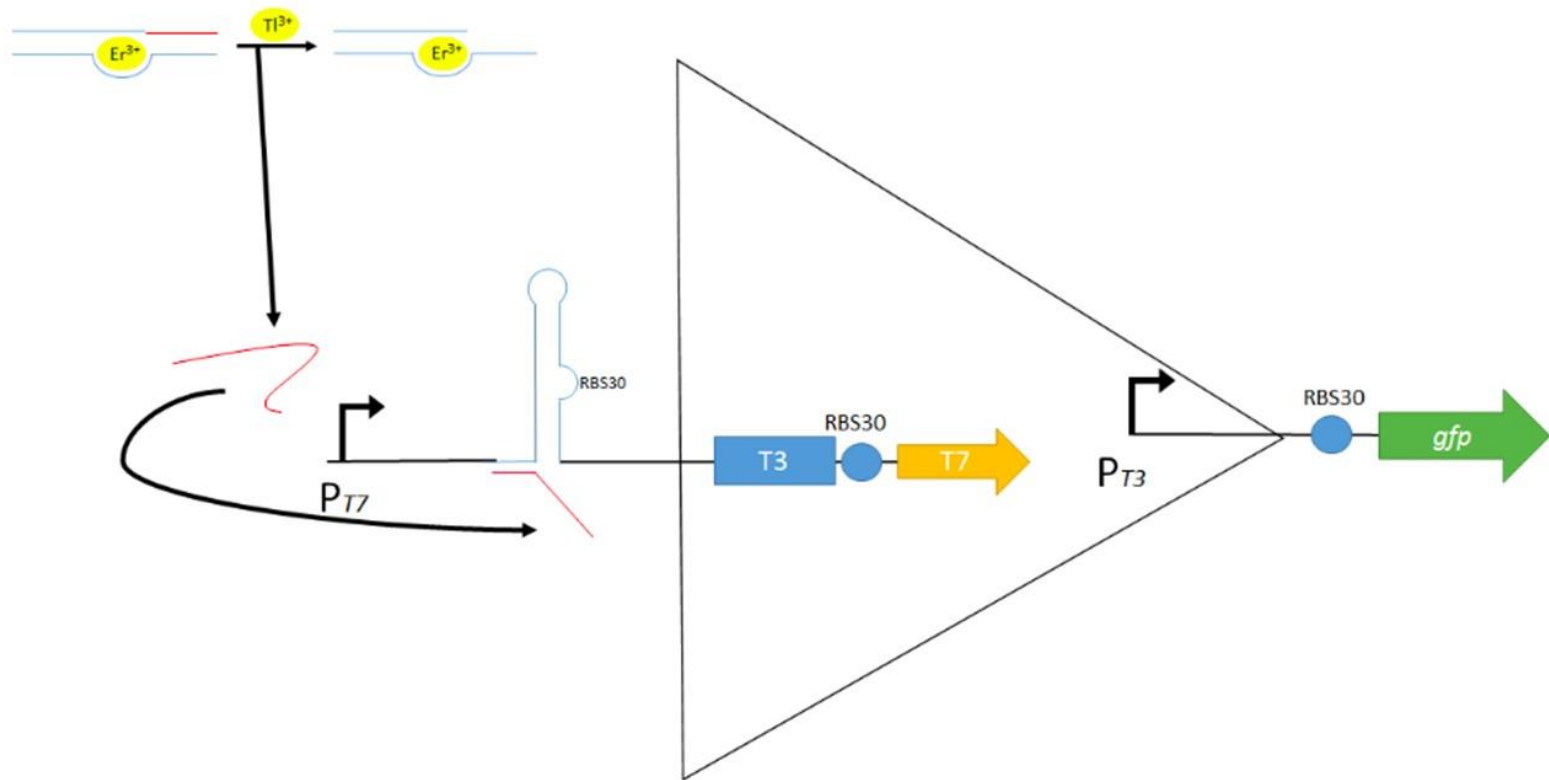
Reporter



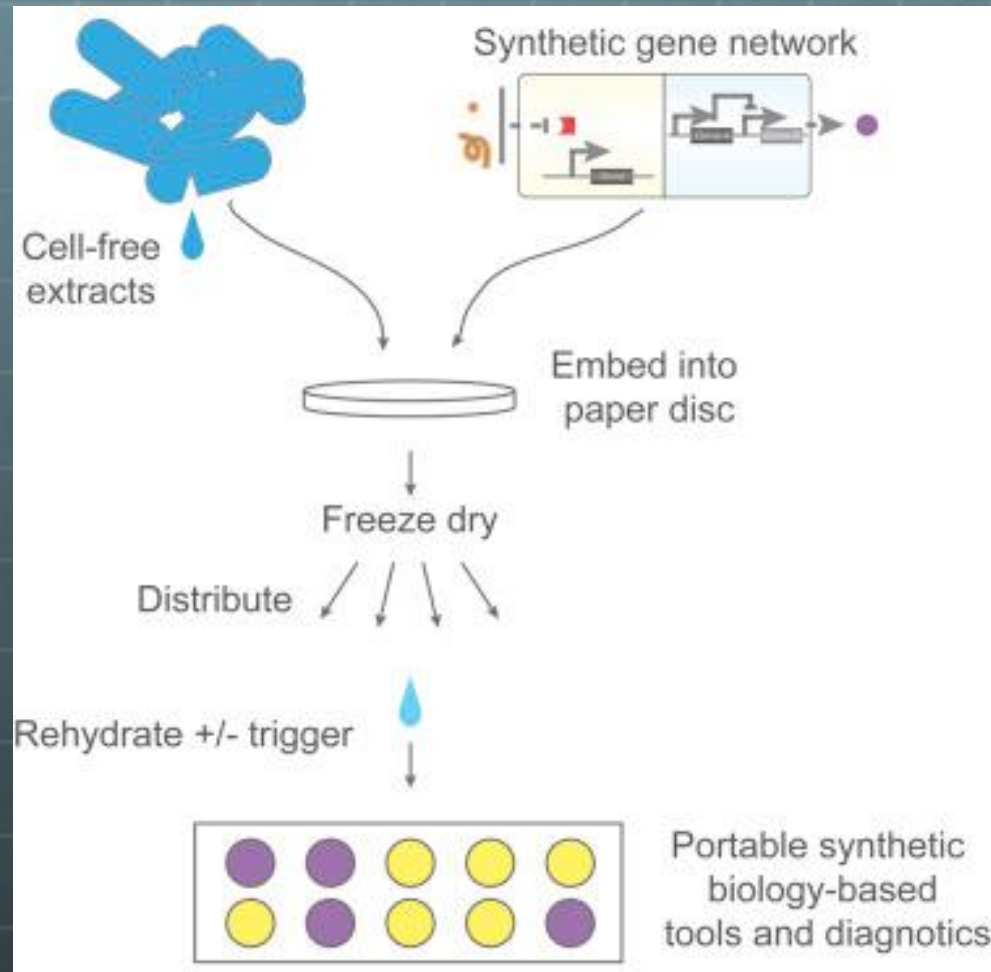
DNAzyme



Amplifier



Cell-Free, Paper-Based Sensor



Paper-Based Synthetic Gene Networks, Pardee et al., 2014

Any Questions?

